2006

Recommended Childhood and Adolescent MDH Immunization Schedule

Range of recommended ages Catch-up Vaccination Preadolescent assessment

Age	Birth	1	2	4	6	12	15	18	24	4-6	11-12	13-14	15	16-18
Vaccine ▼	Dirtii	month	months	months	months	months	months	months	months	years	years	years	years	years
Hepatitis B ¹	НерВ		ſ								HepB S	eries		
Inactivated Polio			P	edia	arix	TM				IPV				
Diphtheria, Tetanus, Pertussis ²			#1	#2	#3		IHIB	I T TM		DTaP	Tdap*		Tdap*	
Haemophilus Influenzae Type b ³			Ped #1	vaxl #2	IIB™	-{ K	JUID	'''						
Measles, Mumps, Rubella						ProQ	uad®			MMR		MN	IR	
Varicella ⁵											Varice	ella		
Meningococcal ⁶								ccines within In line are for	MF	PSV4	MCV4		MCV4	
Pneumococcal ⁷			PCV	PCV	PCV	P		populations	PCV		PP\	/		
Influenza ⁸						 <mark>Influenz</mark> a	 a (yearly)			ln.	fluenza	(yearly)		
Hepatitis A ⁹									HepA Series *					

^{*}MDH clinics will use BOOSTRIX® for adolescent Tdap and VAQTA® for Hep A for infants/young children Brand names are used for simplification and clarification;
Modifications as per ACIP, AAP, AAFP

- 1. Hepatitis B vaccine (HepB). AT BIRTH: All newborns should receive monovalent HepB soon after birth and before hospital discharge. Infants born to mothers who are HBsAg-positive should receive HepB and 0.5mL of hepatitis B immune globulin (HBIG) within 12 hours of birth. Infants born to mothers whose HBsAg status is unknown should receive HepB within 12 hours of birth. The mother should have blood drawn as soon as possible to determine her HBsAg status; if HBsAg-positive, the infant should receive HBIG as soon as possible (no later than age 1 week). For infants born to HBsAg-negative mothers, the birth dose can be delayed in rare circumstances but only if a physician's order to withhold the vaccine and a copy of the mother's original HBsAg-negative laboratory report are documented in the infant's medical record. FOLLOWING THE BIRTH DOSE: The HepB series should be completed with either monovalent HepB or a combination vaccine containing HepB. The second dose should be administered at age 1-2 months. The final dose should be administered at age \$ 24 weeks. It is permissible to administer 4 doses of HepB (e.g., when combination vaccines are given after the birth dose); however, if monovalent HepB is used, a dose at age 4 months is not needed. Infants born to HBsAq-positive mothers should be tested for HBsAq and antibody to HBsAq after completion of the HepB series, at age 9-18 months (generally at the next well-child visit after completion of the vaccine series).
- 2. Diphtheria and tetanus toxoids and acellular pertussis vaccine (DTaP). The fourth dose of DTaP may be administered as early as age 12 months, provided 6 months have elapsed since the third dose and the child is unlikely to return at age 15–18 months. The final dose in the series should be given at age \$4 years. Tetanus and diphtheria toxoids and acellular pertussis vaccine (Tdap adolescent preparation) is recommended at age11–12 years for those who have completed the recommended childhood DTP/DTaP vaccination series and have not received a Td booster dose. Adolescents 13–18 years who missed the 11–12-year Td/Tdap booster dose should also receive a single dose of Tdap if they have completed the recommended childhood DTP/DTaP vaccination series. Subsequent tetanus and diphtheria toxoids (Td) are recommended every10 years.
- 3. Haemophilus influenzae type b conjugate vaccine (Hib). Three Hib conjugate vaccines are licensed for infant use. If PRP-OMP (PedvaxHIB® or ComVax® [Merck]) is administered at ages 2 and 4 months, a dose at age 6 months is not required. DTaP/Hib combination products should not be used for primary immunization in infants at ages 2, 4 or 6 months but can be used as boosters after any Hib vaccine. The final dose in the series should be administered at age \$12 months.
- **4. Measles, mumps, and rubella vaccine (MMR)**. The second dose of MMR is recommended routinely at age 4–6 years but may be administered during any visit, provided at least 4 weeks have elapsed since the first dose and both doses are administered beginning at or after age 12 months. Those who have not previously received the second dose should complete the schedule by age 11–12 years.

- **5. Varicella vaccine**. Varicella vaccine is recommended at any visit at or after age 12 months for susceptible children (i.e., those who lack a reliable history of chickenpox). Susceptible persons aged \$13 years should receive 2 doses administered at least 4 weeks apart.
- **6. Meningococcal vaccine (MCV4)**. Meningococcal conjugate vaccine (MCV4) should be given to all children at the 11–12 year old visit as well as to unvaccinated adolescents at high school entry (15 years of age). Other adolescents who wish to decrease their risk for meningococcal disease may also be vaccinated. All college freshmen living in dormitories should also be vaccinated, preferably with MCV4, although **meningococcal polysaccharide vaccine (MPSV4)** is an acceptable alternative. Vaccination against invasive meningococcal disease is recommended for children and adolescents aged \$2 years with terminal complement deficiencies or anatomic or functional asplenia and certain other high risk groups (see MMWR 2005;54 [RR-7]:1-21); use MPSV4 for children aged 2–10 years and MCV4 for older children, although MPSV4 is an acceptable alternative.
- **7.** Pneumococcal vaccine. The heptavalent pneumococcal conjugate vaccine (PCV) is recommended for all children aged 2–23 months and for certain children aged 24–59 months. The final dose in the series should be given at age \$12 months. Pneumococcal polysaccharide vaccine (PPV) is recommended in addition to PCV for certain high-risk groups. See MMWR 2000; 49(RR-9):1-35.
- 8. Influenza vaccine. Influenza vaccine is recommended annually for children aged \$6 months with certain risk factors (including, but not limited to, asthma, cardiac disease, sickle cell disease, human immunodeficiency virus [HIV], diabetes, and conditions that can compromise respiratory function or handling of respiratory secretions or that can increase the risk for aspiration), healthcare workers, and other persons (including household members) in close contact with persons in groups at high risk (see MMWR 2005;54[RR-8]:1-55). In addition, healthy children aged 6-23 months and close contacts of healthy children aged 0-5 months are recommended to receive influenza vaccine because children in this age group are at substantially increased risk for influenza-related hospitalizations. For healthy persons aged 5-49 years, the intranasally administered, live, attenuated influenza vaccine (LAIV) is an acceptable alternative to the intramuscular trivalent inactivated influenza vaccine (TIV). See MMWR 2005;54(RR-8):1-55. Children receiving TIV should be administered a dosage appropriate for their age (0.25 mL if aged 6-35 months or 0.5 mL if aged \$3 years). Children aged #8 years who are receiving influenza vaccine for the first time should receive 2 doses (separated by at least 4 weeks for TIV and at least 6 weeks for LAIV).
- **9. Hepatitis A vaccine (HepA).** HepA is recommended for all children at 1 year of age (i.e.,12–23 months). The 2 doses in the series should be administered at least 6 months apart. States, counties, and communities with existing HepA vaccination programs for children 2–18 years of age are encouraged to maintain these programs. In these areas, new efforts focused on routine vaccination of 1-year-old children should enhance, not replace, ongoing programs directed at a broader population of children. HepA is also recommended for certain high risk groups (see MMWR 1999; 48[RR-12]1-37).